



# International Filter Products

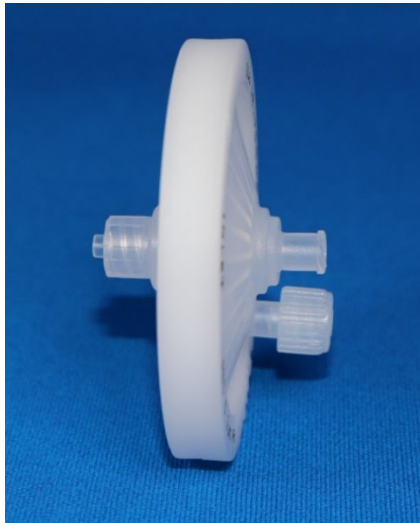
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## STERILE SYRINGE FILTER

These filters are designed for applications where sterility of the affluent is mandatory (Sterile Compounding and Pharmaceutical use). The product contains a pharmaceutical-grade PTFE membrane (Sterilizing Grade) and the filters are built in a 10,000-class clean room, flushed, dried and 100% integrity tested prior to packaging. The product is factory sterilized by Ethylene Oxide gas (ETO) ensuring sterility.

### Technical Data Sheet

**Catalog Number: D65RF100LFLM-PH-ETO**



#### **Materials of Construction**

**Filter Membrane:** 1.0 µm Pharmaceutical Grade Teflon (PTFE) Sterilizing Grade - Hydrophobic

**Supports:** Polypropylene

**Capsule Body:** Polypropylene

#### **Filter Dimensions and Specifications**

**Outer Diameter:** 73 mm (2.87")

**Capsule Body Thickness:** 10.1 mm (0.397")

**Length, Fitting to Fitting:** 40 mm (1.575")

**Filter Area:** 26 cm<sup>2</sup> (4 in<sup>2</sup>)

**Sterilization:** The filter is factory sterilized by Ethylene Oxide Gas (ETO)

**Inlet Fitting:** Luer Loc Female

**Outlet Fitting:** Luer Loc Male

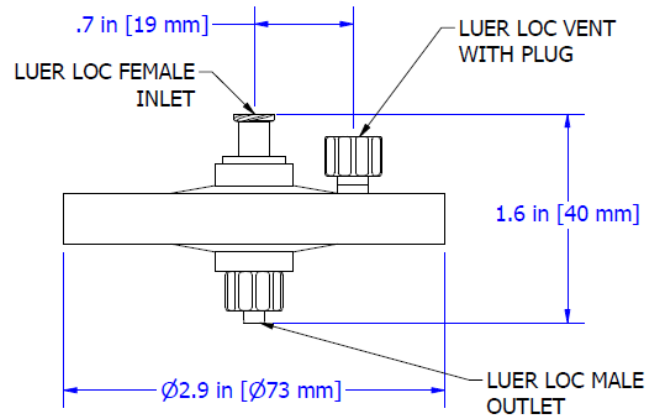
**Volume Filtered:** 460 ml to 2,000 ml (30% - 50% less for Oils)

**Allowable Flowrate:** 34 ml per min to 57 ml per min



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## Operating Parameters

Maximum Working Pressure: 5.5 bar (80 Psi) @ 25 °C (77 °F)

Maximum differential Pressure (forward): 4.1 bar (60 Psi) @ 25 °C (77 °F)

Maximum differential Pressure (reverse): 2.1 bar (30 Psi) @ 25 °C (77 °F)

Maximum Operating Temperature: 80 °C (176 °F)

Recommended Replacement Pressure: 2.4 bar (35.3 Psi)

## Safety

- UPS Bacteria Endotoxins:  $\leq 25$  EU/ml
- Factory Sterilized: Ethylene Oxide Gas (ETO)
- Product was 100% Integrity Tested and flushed with filtered water @ 0.05  $\mu$ m

## Regulatory

- Complies with USP 797 Guidelines
- FDA 21 CFR 177.1655
- USP Class VI - Biological Reactivity
- ISO 10993-Part 1. 5 - Cytotoxicity
- ISO 14001; ISO 13485; OHSMS 18001 Certified.
- ASTM Hemolysis testing
- Human/Veterinarian Use – Applies when the requirements from CGMP CFR part 210 and 211 and additional requirements 21 CFR part 600 and 21 CFR part 680 are used in the aseptic processing.